0			2002/05/2 7 14:04	USPAT; US-PGPUB; EPO; DERWENT	1 same conjugate	6	L4	BRS	4
0			2002/05/2 7 14:04	USPAT; US-PGPUB; EPO; DERWENT	1 same 2	14	L3	BRS	ω
0			2002/05/2 7 13:59	USPAT; US-PGPUB; EPO; DERWENT	25206liposome	25206	L2	BRS	Ν
0			2002/05/2 7 13:58	USPAT; US-PGPUB; EPO; DERWENT	(glucagon-like adj peptide) or glp-1 or glp-2	517	Ľ	BRS	Ъ
Er ro	Error Defin ition	Comm	Time Stamp	DBs	Search Text	Type L # Hits	# 1	Туре	

0			2002/05/2 7 20:15	USPAT; US-PGPUB; EPO; DERWENT	1 same fragment		L3 100	BRS	. ω
0			2002/05/2 7 20:15	JB; IRWENT	1 same analog	182	L2	BRS	2
0			2002/05/2 7 20:14	USPAT; US-PGPUB; EPO; DERWENT	(glucagon-like adj peptide) or glp-1 or glp-2	517	Ľ1	BRS	<u> </u>
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Met-Asn-Thr-OH Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-H-His-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Lys-Tyr-M,:3482.80 [16941-32-5] H-6790.1000

Asn-Thr-NH<sub>2</sub> Asp-Ser-Arg-Arg-Ala-Gin-Asp-Phe-Val-Gin-Trp-Leu-Met-H-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Glu-Tyr-Ser-Lys-Tyr-Leu-(Des-His¹,Gluº) -Glucagon (1-29) amide (human, bovine, porcine) H-2754.0500 H-2754.1000

0.5

g g

135.-

-15 °C

Glucagon antagonist. C148H221N41O47S M,:3358.70 [110084-95-2]

Lit. C.G.Unson et al., Peptides 10, 1171 (1989)

.15 °C Met-Asn-Thr-Lys-Arg-Asn-Lys-Asn-Asn-Ile-Ala-OH Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Glucagon (1-37) (porcine) H-His-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Lys-Tyr-H-6880.0500 H-6880.1000

0.5 mg

430.-785.-

1 mg

Lit. D.Bataille et al., Peptides 2 (suppl. 2), 41 (1981)/ D.Bataille et al., Ann. N.Y. Acad. Sci. 527 (Oxyntomodulin (porcine))

-15 °C Glucagon (19-29) (human, bovine, porcine) H-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-OH H-2758.0005 H-2758.0001

1 mg

35. 145.

This glucagon fragment inhibited both the Ca<sup>2</sup>+ activated and Mg<sup>2</sup>+ dependent ATPase activity C1H89N15O18S and Ca<sup>2</sup>+ transport in liver plasma membranes with an efficiency 1000-fold higher than that of glucagon. It is likely to be the active peptide involved in the inhibition of the liver  $Ca^2+$  pump Lit. A.Mallat et al., Nature **325**, 620 (1987) M;:1352.53 [64790-15-4]

:15°C (GLP-1 amide (human); Preproglucagon (72-107) Glucagon-Like Peptide 1 amide (human) Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-NH<sub>2</sub> Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-H-His-Asp-Glu-Phe-Glu-Arg-His-Ala-Glu-Gly-Thr-Phe-H-6025.1000 H-6025.0500 0.5 mg 1 mg

215.-395.-

.15 ℃

C184H273N51O57 amide (human)) M,:4111.50 [99658-04-5]

Lit. D.J.Drucker et al., Proc. Natl. Acad. Sci. USA **84**, 3434 (1987)/ J.J.Holst et al., FEBS Lett. **211**, 169 (1987)

H-6795.0500 0.5 mg 180. H-6795.1000 1 mg 320.	Glucagon-Like Peptide 1 (7-36) amide H (human) H-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr- Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-lle-Ala-Trp-Leu- Val-Lys-Gly-Arg-NH <sub>2</sub>
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180.-320.-

effect. It is presently considered as the most important incretin hormone. Its action is mediated This GLP-1 fragment is secreted from the lower small intestine and shows a strong insulinotropic receptors expressed by the endocrine pancreatic B-cells. C149H226N40O45 107) amide (human)) M;:3297.68 [107444-51-9] n); Freprogiucagon (/a-

and J.H.Nielsen, FEBS Lett. **229**, 175 (1988) / J.-P.Raufman et al., J. Biol. Chem. **267**, 21432 (1992) / P.A.Martin and A.Faulkner, Comp. Biochem. Biophys. **105A**, 705 (1993) / H.C.Fehmann et al., Lit. B.Kreymann et al., Lancet 1987 II, 1300/ G.I.Bell et al., Nature 304, 368 (1983) / C.Orskov Peptides 15, 453 (1994) / M.A.Nauck et al., Exp. Clin. Endocrinol. Diabetes 105, 187 (1997)

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new (Ser<sup>8</sup>)-Glucagon-Like Peptide 1 (7-36) amide (human) H-4592.1000 H-4592.0500 0.5 mg 1 mg

180.-320.-

Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-NH2 H-His-Ser-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-

((Ser<sup>8</sup>)-GLP-1 (7-36) amide (human); (Ser<sup>79</sup>)-Prepro-

amide against DPP IV without impairing its insulinatropic activity. This may indicate that this modification could improve the potential of GLP-1 in the treatment of type-II diabetes. glucagon (78-107) amide (human))
The replacement of alanine by serine significantly improved the plasma stability of GLP-1 (7-36)

new Glucagon-Like Peptide 2 (human) H-His-Ala-Asp-Gly-Ser-Phe-Ser-Asp-Glu-Met-Asn-Thr-Ile-

C171H266N48O56S (GLP-2 (human); Preproglucagon (126-159) (human)) lle-Gln-Thr-Lys-lle-Thr-Asp-Arg-OH Leu-Asp-Asn-Leu-Ala-Ala-Arg-Asp-Phe-lle-Asn-Trp-Leu-H-4766.0500 H-4766.0001

1 mg 0.5 mg

255.-145.-

with a broad variety of intestinal diseases characterized by intestinal damage and insufficiency. Lit. D.J.Drucker, Trends Endocrinol. Metab. 10, 153 (1999) / D.J.Drucker et al., J. Parenter. Enteral rodents and humans. Currently GLP-2 is used as a potential therapeutic agent for human subjects Like GLP-1, GLP-2 is secreted from enteroendocrine cells in a nutrient dependent manner in both

D-Gluconyl- Val-Leu-Gly-Lys-NHE C27H52N6O10 [121459-49-2] H-8530.0001 H-8530.0005

25 mg 1 mg 5 mg

30.-115.-

Lit. R.Mayer et al., J. Med. Chem. **34**, 3029 (1991) Plasmodium falciparum merozoites ( $IC_{50} = 900 \mu M$ ). Inhibitor of the Plasmodium falciparum proteinase (Ki 480 µM) and of the erythrocyte invasion by H-8530.0025

-15 °C Polypeptide (human) Glucose- Dependent Insulinotropic

(H-Glu(Cys-βNA)-OH)2  $C_{36}H_{40}N_6O_8S_2$ ((H-γ-Glu-Cys-βNA)<sub>2</sub>) (Disulfide bond) (See Gastric Inhibitory Polypeptides and Fragments Page 415) M,:748.88 H-5645.0500 H-5645.1000 K-1650:0050 K-1650.0250 0.5 n. 1 mg 50 mg 250 mg 70. 285. 215.-395.-

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